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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/575,474	04/12/2006	Hiroko Kojima	062405	3422
	7590 02/26/200 I, HATTORI, DANIEL	EXAMINER		
1250 CONNEC	TICUT AVENUE, NV	SHEN, WU CHENG WINSTON		
SUITE 700 WASHINGTOI	N, DC 20036		ART UNIT	PAPER NUMBER
			1632	
			MAIL DATE	DELIVERY MODE
			02/26/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

## Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)	
10/575,474	KOJIMA ET AL.	
Examiner	Art Unit	
WU-CHENG Winston SHEN	1632	

\	WU-CHENG Winston SHEN	1632					
The MAILING DATE of this communication appear	rs on the cover sheet with the c	correspondence add	ress				
THE REPLY FILED 25 January 2008 FAILS TO PLACE THIS APPLICATION IN CONDITION FOR ALLOWANCE.							
1. The reply was filed after a final rejection, but prior to or on the application, applicant must timely file one of the following reapplication in condition for allowance; (2) a Notice of Appea for Continued Examination (RCE) in compliance with 37 CF periods:	plies: (1) an amendment, affidavi I (with appeal fee) in compliance	t, or other evidence, w with 37 CFR 41.31; or	hich places the (3) a Request				
a) The period for reply expires <u>3</u> months from the mailing date of	f the final rejection.						
b) The period for reply expires on: (1) the mailing date of this Adv no event, however, will the statutory period for reply expire late	er than SIX MONTHS from the mailing	g date of the final rejection	n.				
Examiner Note: If box 1 is checked, check either box (a) or (b) MONTHS OF THE FINAL REJECTION. See MPEP 706.07(f).							
Extensions of time may be obtained under 37 CFR 1.136(a). The date or have been filed is the date for purposes of determining the period of exter under 37 CFR 1.17(a) is calculated from: (1) the expiration date of the she set forth in (b) above, if checked. Any reply received by the Office later th may reduce any earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	nsion and the corresponding amount of ortened statutory period for reply origi	of the fee. The appropria nally set in the final Office	ate extension fee e action; or (2) as				
2. The Notice of Appeal was filed on A brief in complia filing the Notice of Appeal (37 CFR 41.37(a)), or any extens	ion thereof (37 CFR 41.37(e)), to	avoid dismissal of the	s of the date of e appeal. Since a				
Notice of Appeal has been filed, any reply must be filed with AMENDMENTS	nin the time period set forth in 37 (	CFR 41.37(a).					
	t prior to the date of filing a brief	will not be entered be	cause				
<ul> <li>3.  The proposed amendment(s) filed after a final rejection, but prior to the date of filing a brief, will not be entered because <ul> <li>(a) They raise new issues that would require further consideration and/or search (see NOTE below);</li> <li>(b) They raise the issue of new matter (see NOTE below);</li> <li>(c) They are not deemed to place the application in better form for appeal by materially reducing or simplifying the issues for</li> </ul> </li> </ul>							
appeal; and/or	Tioninion appear by materially rec	adding or simplifying the	ie issues ioi				
(d) ☐ They present additional claims without canceling a co	rresponding number of finally reje	ected claims.					
NOTE: (See 37 CFR 1.116 and 41.33(a)).							
4. The amendments are not in compliance with 37 CFR 1.121 5. Applicant's reply has overcome the following rejection(s): <u>I</u> by Kumta et al., is withdrawn in light of Applicant's arguments, wh	he rejection of claims 9-12 under	•	•				
6. Newly proposed or amended claim(s) would be allow non-allowable claim(s).	wable if submitted in a separate, t	imely filed amendmer	nt canceling the				
7. For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is provided the status of the claim(s) is (or will be) as follows:		l be entered and an ex	xplanation of				
Claim(s) allowed: Claim(s) objected to:							
Claim(s) objected to:  Claim(s) rejected: <u>9-12</u> .  Claim(s) withdrawn from consideration:							
AFFIDAVIT OR OTHER EVIDENCE							
8. The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good and swas not earlier presented. See 37 CFR 1.116(e).							
9. The affidavit or other evidence filed after the date of filing a entered because the affidavit or other evidence failed to ove showing a good and sufficient reasons why it is necessary a	ercome <u>all</u> rejections under appea	al and/or appellant fails	s to provide a				
10.   The affidavit or other evidence is entered. An explanation							
REQUEST FOR RECONSIDERATION/OTHER							
11. The request for reconsideration has been considered but on See Continuation Sheet.		condition for allowan	ce because:				
<ul><li>12. ☐ Note the attached Information <i>Disclosure Statement</i>(s). (P</li><li>13. ☐ Other:</li></ul>	I O/SB/08) Paper No(s)						
	Malaria Dantariia						
	Nalarie Bertoglio/ Primary Examiner Art Unit 1632						

Continuation of 11. does NOT place the application in condition for allowance because:

Applicant's arguments have failed to overcome the rejection of claims 9-12 under 35 U.S.C. 102(e) and under 35 U.S.C. 102(a) as being anticipated by Doll et al. (Doll et al., U.S. Patent application Publication 2003/0235564, publication date Dec. 25, 2003, filed on May 13, 2003).

Applicant argues that Doll does not inherently disclose an implant including the claimed adsorption. Doll does not explicitly describe how Runx2 is incorporated into a porous material. Applicant also argues that specific procedures must be undertaken in order to ensure adsorption. Applicant further cites the disclosure on pages 7, 11, and 16, which indicates that a degasified condition under a reduced pressure of 100 to 150 mmHg for 3 hours or longer is required for the claimed adsorption to occur.

In response, the Examiner notes that claim 9 is a product claim and reads as follows: "An implant consisting of a bioadaptable porous material on which an adenoviral or retroviral vector carrying a gene encoding an osteo-inducible transcription factor Cbfal is adsorbed." Applicant's arguments focus on the process how the implant being made requires a specified condition. However, the Examiner notes that the degasified reduced pressure condition is not recited in the claim. The Examiner maintains the position that Doll et al. anticipates the claimed invention because Doll et al. disclose beta-TCP, a bio-compatible biodegradable polymeric matrix, in the context of a pharmacetical composition comprising a polynucleotide encoding Cbfa1, in a pharmaceutically accetable carrier. It is noted that a pharmaceutically accetable carrier reads on water. Considering a pharmaceutical composition comprises beta-TCP and a polynucleotide encoding Cbfa1 dissolved in water, the claimed adsorption process will inherently occur. Therefore, Doll et al. anticipate the cliamed invention. Further information regarding product-by-process claims is provided below.

"Where, as here, the claimed and prior art products are identical or substantially identical, or are produced by identical or substantially identical processes, the PTO can require an applicant to prove that the prior art products do not necessarily or inherently possess the characteristics of his claimed product (In re Ludtke). Whether the rejection is based on "inherency" under 35 USC 102, on "prima facie obviousness" under 35 USC 103, jointly or alternatively, the burden of proof is the same, and its fairness is evidenced by the PTO's inability to manufacture products or to obtain and compare prior art products. In re Best, Bolton, and Shaw, 195 USPQ 430, 433 (CCPA 1977) citing In re Brown, 59 CCPA 1036, 459 F.2d 531, 173 USPQ 685 (1972))."

"When the PTO shows a sound basis for believing that the products of the applicant and the prior art are the same, the applicant has the burden of showing that they are not." In re Spada, 911 F.2d 705, 709, 15 USPQ2d 1655, 1658 (Fed. Cir. 1990). Therefore, the prima facie case can be rebutted by evidence showing that the prior art products do not necessarily possess the characteristics of the claimed product. In re Best, 562 F.2d at 1255, 195 USPQ at 433. See also Titanium Metals Corp. v. Banner, 778 F.2d 775, 227 USPQ 773 (Fed. Cir. 1985), In re Ludtke, 441 F.2d 660, 169 USPQ 563 (CCPA 1971), Northam Warren Corp. v. D. F. Newfield Co., 7 F. Supp. 773, 22 USPQ 313 (E.D.N.Y. 1934.) See MPEP 2113 and MPEP 2112.01.

Even though product-by-process claims are limited by and defined by the process; determination of patentability is based on the product itself. The patentability of a product does not depend on its method of production. If the product in the product-by-process claim is the same as or obvious from a product of the prior art, the claim is unpatentable even though the prior product was made by a different process." In re Thorpe, 777 F.2d 695, 698, 227 USPQ 964, 966 (Fed. Cir. 1985).

"The Patent Office bears a lesser burden of proof in making out a case of prima facie obviousness for product-by-process claims because of their peculiar nature" than when a product is claimed in the conventional fashion. In re Fessmann, 489 F.2d 742, 744, 180 USPQ 324, 326 (CCPA 1974). Once the examiner provides a rationale tending to show that the claimed product appears to be the same or similar to that of the prior art, although produced by a different process, the burden shifts to applicant to come forward with evidence establishing an unobvious difference between the claimed product and the prior art product. In re Marosi, 710 F.2d 798, 802, 218 USPQ 289, 292 (Fed. Cir. 1983).

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